

# SCORE Search Results Details for Application 10552515 and Search Result 20080624\_135827\_us-10-552-515-1\_copy\_157\_933.rag.

<a href="#">Score Home</a>	<a href="#">Retrieve Application</a>	<a href="#">SCORE System</a>	<a href="#">SCORE</a>	<a href="#">Comments /</a>
<a href="#">Page</a>	<a href="#">List</a>	<a href="#">Overview</a>	<a href="#">FAQ</a>	<a href="#">Suggestions</a>

This page gives you Search Results detail for the Application 10552515 and Search Result 20080624\_135827\_us-10-552-515-1\_copy\_157\_933.rag.

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GenCore version 6.2.1  
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OM protein - protein search, using sw model

Run on: June 24, 2008, 15:17:27 ; Search time 267 Seconds  
(without alignments)  
1751.538 Million cell updates/sec

Title: US-10-552-515-1\_COPY\_157\_933  
Perfect score: 4123  
Sequence: 1 QQDVQDGNTTVHYALLSASW.....SELSSHWTPTVTPKASQLQQ 777

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 3405708 seqs, 601879884 residues

Total number of hits satisfying chosen parameters: 3405708

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : A\_Geneseq\_200711:\*  
1: geneseqp1980s:\*  
2: geneseqp1990s:\*  
3: geneseqp2000:\*  
4: geneseqp2001:\*  
5: geneseqp2002:\*  
6: geneseqp2003a:\*  
7: geneseqp2003b:\*

8: geneseqp2004a:\*  
 9: geneseqp2004b:\*  
 10: geneseqp2005:\*  
 11: geneseqp2006:\*  
 12: geneseqp2007:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	% Query		DB	ID	Description
		Match	Length			
1	4123	100.0	933	8	ADT77664	Adt77664 Splice va
2	4123	100.0	933	11	AEL84788	Ael84788 Tumor mar
3	3739	90.7	885	10	AEB13426	Aeb13426 Human pro
4	3572	86.6	843	10	AEB13424	Aeb13424 Human pro
5	3031.5	73.5	898	4	ABG15488	Abg15488 Novel hum
6	1502.5	36.4	920	6	ADB64420	Adb64420 Human pro
7	1482.5	36.0	920	6	ABP58666	Abp58666 Human dih
8	1467.5	35.6	981	8	ADK52114	Adk52114 Human ato
9	1467.5	35.6	981	12	AEN06206	Aen06206 Human eso
10	1461.5	35.4	840	11	AEG11146	Aeg11146 Human tra
11	1456.5	35.3	960	11	AEG11142	Aeg11142 Human tra
12	1452.5	35.2	1017	12	AFB77190	Afb77190 Mouse TM-
13	1437	34.9	1003	7	ADG48280	Adg48280 Human ret
14	1412.5	34.3	913	11	AEH82071	Aeh82071 Human gna
15	1378.5	33.4	1219	4	ABB62812	Abb62812 Drosophil
16	1378.5	33.4	1219	10	AFB95185	Afb95185 Fruit fly
17	1369	33.2	910	6	ADC42854	Adc42854 REMAP pro
18	1369	33.2	910	11	AEL84658	Ael84658 Tumor mar
19	1367.5	33.2	712	11	AEG11145	Aeg11145 Human tra
20	1344	32.6	1075	4	ABB65993	Abb65993 Drosophil
21	1344	32.6	1075	10	AFC04729	Afc04729 Fruit fly
22	1159.5	28.1	1058	4	ABB65022	Abb65022 Drosophil
23	1159.5	28.1	1058	10	AFC01816	Afc01816 Fruit fly
24	1154	28.0	596	6	ADB64387	Adb64387 Human pro
25	1061.5	25.7	594	4	AAB92637	Aab92637 Human pro
26	1061.5	25.7	594	5	ABP43811	Abp43811 FLJ10261
27	1061.5	25.7	594	8	ADJ75429	Adj75429 Marker ge
28	1061.5	25.7	594	8	ADN04848	Adn04848 Antipsori
29	1061.5	25.7	594	11	AEG11143	Aeg11143 Human FLJ
30	1024.5	24.8	782	6	ADX42387	Adx42387 Human col
31	1024.5	24.8	782	7	ADT95905	Adt95905 Colon can
32	1024.5	24.8	782	8	ADQ96288	Adq96288 T cell ac
33	1024.5	24.8	782	8	ADQ96104	Adq96104 T cell ac
34	912.5	22.1	475	6	ADB64962	Adb64962 Human pro

35	873.5	21.2	642	7	ADM05798	Adm05798 Human pro
36	873.5	21.2	642	10	AEC88728	Aec88728 Human cDN
37	873.5	21.2	642	11	AEG11144	Aeg11144 Human FLJ
38	819.5	19.9	443	5	ABP41785	Abp41785 Human ova
39	784.5	19.0	390	5	ABB90382	Abb90382 Human pol
40	735	17.8	139	5	AAE24066	Aae24066 Human pro
41	722.5	17.5	360	4	AAM40391	Aam40391 Human pol
42	711.5	17.3	346	8	ADP29628	Adp29628 Human sec
43	695.5	16.9	608	8	ADQ96298	Adq96298 T cell ac
44	695.5	16.9	608	8	ADQ96286	Adq96286 T cell ac
45	684.5	16.6	483	7	ADM05305	Adm05305 Human pro

## ALIGNMENTS

## RESULT 1

ADT77664

ID ADT77664 standard; protein; 933 AA.

XX

AC ADT77664;

XX

DT 15-JUN-2007 (revised)

DT 13-JAN-2005 (first entry)

XX

DE Splice variant-novel gene expressed in prostate (SV-NGEP) polypeptide.

XX

KW Splice variant-novel gene expressed in prostate; SV-NGEP; human;  
 KW prostate cancer; cytostatic; gene therapy; immunotherapy; BOND\_PC;  
 KW NGEP long variant; NGEP long variant [Homo sapiens]; G05886.

XX

OS Homo sapiens.

XX

FH Key Location/Qualifiers

FT Domain 1. .345

FT /label= Cytoplasmic

FT Region 157. .933

FT /note= "An immunogenic fragment comprising 8 consecutive  
 FT amino acids that specifically binds to an antibody that  
 FT specifixally binds to a polypeptide comprising amino  
 FT acids 157-933 is referred to in Claim 1"

FT Region 170. .178

FT /note= "Epitope, predicted to bind HLA2-01"

FT Region 215. .223

FT /note= "Epitope, predicted to bind HLA2-01"

FT Region 258. .266

FT /note= "Epitope, predicted to bind HLA2-01"

FT Domain 346. .368

FT /label= Transmembrane

FT	Domain	369. .421
FT		/label= External
FT		/note= "Cell surface"
FT	Region	403. .411
FT		/note= "Epitope, predicted to bind HLA2-01"
FT	Domain	422. .441
FT		/label= Transmembrane
FT	Region	427. .435
FT		/note= "Epitope, predicted to bind HLA2-01"
FT	Domain	442. .501
FT		/label= Cytoplasmic
FT	Domain	502. .524
FT		/label= Transmembrane
FT	Domain	525. .543
FT		/label= External
FT		/note= "Cell surface"
FT	Domain	544. .566
FT		/label= Transmembrane
FT	Region	557. .565
FT		/note= "Epitope, predicted to bind HLA2-01"
FT	Region	562. .570
FT		/note= "Epitope, predicted to bind HLA2-01"
FT	Domain	567. .586
FT		/label= Cytoplasmic
FT	Domain	587. .609
FT		/label= Transmembrane
FT	Domain	610. .714
FT		/label= External
FT		/note= "Cell surface"
FT	Domain	715. .737
FT		/label= Transmembrane
FT	Domain	738. .761
FT		/label= Cytoplasmic
FT	Domain	762. .784
FT		/label= Transmembrane
FT	Domain	785. .933
FT		/label= External
FT		/note= "Cell surface"
FT	Region	846. .854
FT		/note= "Epitope, predicted to bind HLA2-01"

XX

PN W02004092213-A1.

XX

PD 28-OCT-2004.

XX

PF 05-APR-2004; 2004WO-US010588.

XX

PR 08-APR-2003; 2003US-0461399P.

XX

PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.

XX

PI Pastan I, Bera TK, Lee B;

XX

DR WPI; 2004-758338/74.

DR N-PSDB; ADT77665.

DR PC:NCBI; gi48093524.

XX

PT New Splice Variant-Novel Gene Expressed in Prostate polypeptide or  
PT encoding nucleic acid molecule for diagnosing, preventing or treating  
cancer, especially prostate cancer.

XX

PS Claim 1; SEQ ID NO 1; 88pp; English.

XX

CC The present sequence is the protein sequence of splice variant-novel gene  
CC expressed in prostate (SV-NGEP). SV-NGEP is identical to NGEF from amino  
CC acid 1-157, diverging from amino acid 158. Expression analysis in 76  
CC normal and foetal tissues showed SV-NGEP to be strongly expressed only in  
CC a prostate sample. Claimed methods for detecting prostate cancer in a  
CC subject comprise: contacting the sample with an antibody that  
CC specifically binds a SV-NGEP polypeptide and detecting the formation of  
CC an immune complex; or detecting an increase in expression of SV-NGEP  
CC polypeptide or mRNA. Antibodies to an SV-NGEP polypeptide can be used to  
CC detect metastatic prostate cancer cells at locations other than the  
CC prostate. A claimed method for producing an immune response against a  
CC cell expressing SV-NGEP, for example in a subject with prostate cancer,  
CC comprises administering the polypeptide, or a polynucleotide encoding it,  
CC to produce an immune response that decreases growth of the prostate  
CC cancer. A claimed method for inhibiting the growth of a malignant cell  
CC that expresses SV-NGEP comprises culturing cytotoxic T lymphocytes (CTLs)  
CC with SV-NGEP to produce activated CTLs that recognise an NGEF expressing  
CC cell, and contacting the malignant cell with the activated CTLs.  
CC Alternatively, growth of a malignant cell is inhibited by contact with an  
CC antibody that specifically binds an SV-NGEP polypeptide, where the  
CC antibody is linked to an effector molecule (chemotherapeutic agent or  
CC toxin) that inhibits growth of the malignant cell. This may be performed  
CC in vivo. Kits for detecting an SV-NGEP polypeptide or polynucleotide in a  
CC sample are also claimed.

CC

CC Revised record issued on 15-JUN-2007 : Enhanced with precomputed  
CC information from BOND.

XX

SQ Sequence 933 AA;

Query Match	100.0%;	Score 4123;	DB 8;	Length 933;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches 777;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

Qy 1 QQDVQDGNNTTVHYALLSASWAVLCYYAEDLRKLPLQELPNQASNWSAGLLAWLGIPNVL 60



Db 877 ESVEIKVKREYYLAKQALAENEVLFGTNGTKDEQPKGSELSSHWTPTVTPKASQLQQ 933

## RESULT 2

AEL84788

ID AEL84788 standard; protein; 933 AA.

XX

AC AEL84788;

XX

DT 18-OCT-2007 (revised)

DT 15-JUN-2007 (revised)

DT 28-DEC-2006 (first entry)

XX

DE Tumor marker gene NGEP SEQ ID NO 155.

XX

KW cytostatic; diagnosis; prognosis; tumor marker; gene expression;

KW drug screening; cancer; neoplasm; NGEP; BOND\_PC; NGEP long variant;

KW G05886.

XX

OS Homo sapiens.

XX

PN W02006110593-A2.

XX

PD 19-OCT-2006.

XX

PF 07-APR-2006; 2006WO-US013172.

XX

PR 07-APR-2005; 2005US-0669342P.

PR 11-OCT-2005; 2005US-0725982P.

XX

PA (MACR-) MACROGENICS INC.

XX

PI Von Haller PD, Schummer M, Meyer DW, Schubert LA, Tjoelker LW;

XX

DR WPI; 2006-814687/82.

DR N-PSDB; AEL84787.

DR REFSEQ; NP\_001001891.

DR PC:NCBI; gi48093524.

XX

PT Detecting or diagnosing cancer in a subject comprises determining  
 PT expression of at least one gene, and comparing level of expression to a  
 PT control sample from a normal subject, where increased expression level  
 PT indicates cancer.

XX

PS Claim 8; SEQ ID NO 155; 583pp; English.

XX

CC The invention describes a method of detecting or diagnosing cancer in a  
 CC subject comprising determining the expression level of at least one gene,  
 CC and comparing the level of expression to a corresponding control sample

from a normal subject, where cancer is detected or diagnosed if there is an increase in the expression level of the gene relative to the expression in the control sample. Also described are: identifying a compound to be tested for its ability to prevent, treat, manage, or ameliorate cancer or its symptom; a compound identified by the method; treating cancer in a patient; treating a cancer in a subject that is fully or partially refractory to a first treatment in a patient; and a pharmaceutical composition comprising an amount of an antibody selected from anti-SLC12A2, anti-FLJ23375, anti-GRM5, anti-TAS2R1, anti-NRXN2, anti-C14orf160, anti-MGC 15668, anti-MGC33486, anti-TMEM16F, anti-FAT, anti-KIAA0195, anti-LRFN, anti-NFASC, anti-BAT2D1, anti-MGC2963, anti-KIAA0685, anti-EDG3, anti-GGTL3, anti-PLVAP, anti-FLJ31528, anti-FLJ90709, anti-VEZATIN, anti-TMPRSS9, anti-ATP13A5, anti-PKHD1L1, anti-C2orf18, anti-ANKRD22, anti-FAM62B, anti-LOC57168, anti-CDKAL1, anti-SLC39A3v1, anti-SLC39A3v2, anti-BAT5, anti-TM9SF4, anti-DC2, anti-VAPB, anti-XTPTPB, anti-TACSTD2, anti-FNDC3A, anti-GK001, anti-OCIAD2, anti-PR01855, anti-C20orf3, anti-SDFR1, anti-FLJ20481, anti-LENG4, anti-FLJ12443, anti-ARP5 Long, anti-ARP5 Short, anti-TMD0645, anti-NGEP, anti-IL1RAP1, anti-PLXNB1, anti-ATP2B2, anti-FLJ11848, anti-ENTPD2, anti-PPM1H, anti-KRTKAP3, anti-KCNC3, anti-TM9SF1, anti-ULBP1, anti-C19orf26, anti-KIAA830, anti-KIAA1244, anti-KIAA1797, anti-MGC26856, anti-NETO2, anti-SUSD2, anti-FOLR2, anti-EMR2, ENTPD1, anti-ATP10B, anti-PTK7, anti-FLJ14681, anti-C20orf22, anti-FLJ14281, anti-FAM8A1, anti-TMED7, anti-C20orf108, anti-ATAD1, anti-GPR154, anti-C14orf27, anti-OSAP, anti-FAD104, anti-FLJ90492, anti-SLC27A3, anti-RON, anti-ATP13A1, anti-DKFZP564D166, anti-ESSPL, anti-EXTL3, anti-KAIL, anti-KIAA0960, anti-MTRNL, anti-SLC27A1, anti-GRIA, anti-OR4M1, anti-KIAA1679, or anti-UPK-1b antibody, and a pharmaceutical carrier. The methods are useful for detecting, diagnosing, and treating cancer, e.g. colon, lung, ovary, prostate, pancreas, or bladder cancer. This is the amino acid sequence of NGEP, altered levels of expression are useful in the diagnosis or prognosis of cancer.

Revised record issued on 18-OCT-2007 : Enhanced with precomputed information from BOND.

Sequence 933 AA;

Query Match	100.0%;	Score 4123;	DB 11;	Length 933;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches	777;	Conservative	0;	Mismatches 0; Indels 0; Gaps 0;

  

Qy	1	QQDVQDGNNTTVHYALLSASWAVLCYYAEDRLRLKPLQELPNQASNWSAGLLAWLGIPNVL	60
Db	157	QQDVQDGNNTTVHYALLSASWAVLCYYAEDRLRLKPLQELPNQASNWSAGLLAWLGIPNVL	216
Qy	61	LEVVPDVPPEYYSCRFRVKNLPRFLGSDNQDTFFFTSKRHQILFEILAKTPYGHEKKNLL	120
Db	217	LEVVPDVPPEYYSCRFRVKNLPRFLGSDNQDTFFFTSKRHQILFEILAKTPYGHEKKNLL	276



Qy	121	GIHQLLAEGVLSAAFLPHDGPFTKTPPEGPQAPRLNQRQVLFQHWARWGKWNKYQPLDHVR	180
Db	277	GIHQLLAEGVLSAAFLPHDGPFTKTPPEGPQAPRLNQRQVLFQHWARWGKWNKYQPLDHVR	336
Qy	181	RYFGEKVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	240
Db	337	RYFGEKVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	396
Qy	241	CLDCPFWLLSSACALAQAGRLFHDGGTVFFSLFMALWAVLLLEYWKRKSATLAYRWDCSD	300
Db	397	CLDCPFWLLSSACALAQAGRLFHDGGTVFFSLFMALWAVLLLEYWKRKSATLAYRWDCSD	456
Qy	301	YEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVMCLV	360
Db	457	YEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVMCLV	516
Qy	361	SIILYRAIMAIVVSRSGNTLLAAWASRIASLTGSVVNLVFIILISKIYVSLAHVLTREW	420
Db	517	SIILYRAIMAIVVSRSGNTLLAAWASRIASLTGSVVNLVFIILISKIYVSLAHVLTREW	576
Qy	421	HRTQTKFEDAFTLKVFIFQFVNFSYSSPVYIAFFKGRFVGYPGNYHTLFGVRNEECAAGGC	480
Db	577	HRTQTKFEDAFTLKVFIFQFVNFSYSSPVYIAFFKGRFVGYPGNYHTLFGVRNEECAAGGC	636
Qy	481	LIELAQELLVIMVGKQVINNMQEVLI PKLKGWWQKFR LRSKKRKAGASAGASQGPWEDDY	540
Db	637	LIELAQELLVIMVGKQVINNMQEVLI PKLKGWWQKFR LRSKKRKAGASAGASQGPWEDDY	696
Qy	541	ELVPCEGLFDEYLEMVLQFGFVTIFVAACPLAPLFALLNNWVEIRLDARKFVCEYRRPVA	600
Db	697	ELVPCEGLFDEYLEMVLQFGFVTIFVAACPLAPLFALLNNWVEIRLDARKFVCEYRRPVA	756
Qy	601	ERAQDIGIWPHILAGLTHLAVISNAFLAFSSDFLPRAYRWTRAHDLRGFLNFTLARAP	660
Db	757	ERAQDIGIWPHILAGLTHLAVISNAFLAFSSDFLPRAYRWTRAHDLRGFLNFTLARAP	816
Qy	661	SSFAAAHNRTCryRAFRDDDGHSQTYWNLLAIRLAFVIVFEHVVSFVGRLLDLLVPDIP	720
Db	817	SSFAAAHNRTCryRAFRDDDGHSQTYWNLLAIRLAFVIVFEHVVSFVGRLLDLLVPDIP	876
Qy	721	ESVEIKVKREYYLAKQALAENEVLFGTNGTKDEQPKGSELSSHWPFTVPKASQLQQ	777
Db	877	ESVEIKVKREYYLAKQALAENEVLFGTNGTKDEQPKGSELSSHWPFTVPKASQLQQ	933

RESULT 3

AEB13426

ID AEB13426 standard; protein; 885 AA.

XX  
AC AEB13426;  
XX  
DT 22-SEP-2005 (first entry)  
XX  
DE Human prostate specific polypeptide #2.  
XX  
KW Screening; diagnosis; drug delivery; prostate specific polypeptide;  
KW cancer; prostate tumor; cytostatic; neoplasm.  
XX  
OS Homo sapiens.  
XX  
PN WO2005062788-A2.  
XX  
PD 14-JUL-2005.  
XX  
PF 16-DEC-2004; 2004WO-US042406.  
XX  
PR 22-DEC-2003; 2003US-0531809P.  
XX  
PA (AVAL-) AVALON PHARM INC.  
XX  
PI Weigle B, Ebner R;  
XX  
DR WPI; 2005-497793/50.  
DR N-PSDB; AEB13425.  
XX  
PT Novel isolated prostate specific polypeptide, useful for treating cancer,  
PT and identifying agent that modulates activity of cancer related gene.  
XX  
PS Claim 12; SEQ ID NO 5; 59pp; English.  
XX  
CC The invention relates to an isolated prostate specific polypeptide  
CC comprising one or more immunogenic fragments. The invention also relates  
CC to a method of identifying an agent that modulates the activity of a  
CC cancer related gene involving contacting a compound with a cell  
CC containing a gene under conditions promoting the expression of the gene,  
CC detecting a difference in expression of the gene relative to when the  
CC compound is not present and identifying an agent that modulates the  
CC activity of a cancer related gene, a method of identifying an anti-  
CC neoplastic agent involving contacting a cell exhibiting neoplastic  
CC activity with a compound first identified as a cancer related gene  
CC modulator using and determining a decrease in neoplastic activity after  
CC contacting, when compared to when the contacting does not occur, or  
CC administering an agent first identified to an animal exhibiting a cancer  
CC condition and detecting a decrease in cancerous condition, a method of  
CC determining the cancerous status of a cell involving determining an  
CC increase in the level of expression in a cell of a gene where an elevated  
CC expression relative to a known non-cancerous cell indicates a cancerous

state or potentially cancerous state, an antibody that reacts with a prostate specific polypeptide, an immunoconjugate comprising the antibody and a cytotoxic agent, a method of treating cancer involving contacting a cancerous cell in vivo with an agent having activity against a prostate specific polypeptide and an immunogenic composition the prostate specific polypeptide. The prostate specific polypeptide is useful for identifying an agent that modulates the activity of a cancer related gene. The immunogenic composition is useful for treating cancer, preferably prostate cancer in an animal, e.g. human, which involves administering the immunogenic composition that is sufficient to elicit the production of cytotoxic T lymphocytes specific for the prostate specific polypeptide. The invention is useful for identifying anti-neoplastic agents. This sequence represents a human prostate specific polypeptide of the invention.

XX

SQ Sequence 885 AA;

Query Match 90.7%; Score 3739; DB 10; Length 885;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 702; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy	1	QQDVQDGNNTTVHYALLSASWAVLCYYAEDLRKLPLQELPNQASNWSAGLLAWLGIPNVL	60
Db	158	QQDVQDGNNTTVHYALLSASWAVLCYYAEDLRKLPLQELPNQASNWSAGLLAWLGIPNVL	217
Qy	61	LEVVPDVPPEYYSCRFRVKNLPRFLGSDNQDTFFTSTKRHQILFEILAKTPYGHEKKNLL	120
Db	218	LEVVPDVPPEYYSCRFRVKNLPRFLGSDNQDTFFTSTKRHQILFEILAKTPYGHEKKNLL	277
Qy	121	GIHQLLAEGVLSAAFLPHDGFKTPPEGPQAPRLNQRQVLFQHWARWGKWNKYQPLDHVR	180
Db	278	GIHQLLAEGVLSAAFLPHDGFKTPPEGPQAPRLNQRQVLFQHWARWGKWNKYQPLDHVR	337
Qy	181	RYFGEKVVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	240
Db	338	RYFGEKVVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	397
Qy	241	CLDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAVLLLEYWKRKSATLAYRWDCSD	300
Db	398	CLDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAVLLLEYWKRKSATLAYRWDCSD	457
Qy	301	YEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVMCLV	360
Db	458	YEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVMCLV	517
Qy	361	SIILYRAIMAIVVSRSGNTLLAAWASRIASLTGSVVNLVFIILSKIYVSLAHVLTWRWEM	420
Db	518	SIILYRAIMAIVVSRSGNTLLAAWASRIASLTGSVVNLVFIILSKIYVSLAHVLTWRWEM	577

Qy	421	HRTQTKFEDAFILKVFIFQFVNFISSPVYIAFFKGRFVGYPGNYHTLFGVRNEECAAGGC	480
Db	578	HRTQTKFEDAFILKVFIFQFVNFISSPVYIAFFKGRFVGYPGNYHTLFGVRNEECAAGGC	637
Qy	481	LIELAQELLVIMVGKQVINNMQEVLIPLKLGWWQKFRRLRSKKRKAGASAGASQGPWEDDY	540
Db	638	LIELAQELLVIMVGKQVINNMQEVLIPLKLGWWQKFRRLRSKKRKAGASAGASQGPWEDDY	697
Qy	541	ELVPCEGLFDEYLEMVLQFGFVTIFVAACPLAPLFAALLNNWVEIRLDARKFVCEYRRPVA	600
Db	698	ELVPCEGLFDEYLEMVLQFGFVTIFVAACPLAPLFAALLNNWVEIRLDARKFVCEYRRPVA	757
Qy	601	ERAQDIGIWFHILAGLTHLAVISNAFLAFSSDFLPRAYRWTRAHDLRGFLNFTLARAP	660
Db	758	ERAQDIGIWFHILAGLTHLAVISNAFLAFSSDFLPRAYRWTRAHDLRGFLNFTLARAP	817
Qy	661	SSFAAAHNRTCryRAFRDDDGHSQTYWNLALIRLAFVIVFE	702
Db	818	SSFAAAHNRTCryRAFRDDDGHSQTYWNLALIRLAFVIVFE	859

## RESULT 4

AEB13424

ID AEB13424 standard; protein; 843 AA.

XX

AC AEB13424;

XX

DT 22-SEP-2005 (first entry)

XX

DE Human prostate specific polypeptide #1.

XX

KW Screening; diagnosis; drug delivery; prostate specific polypeptide;  
 KW cancer; prostate tumor; cytostatic; neoplasm.

XX

OS Homo sapiens.

XX

PN W02005062788-A2.

XX

PD 14-JUL-2005.

XX

PF 16-DEC-2004; 2004WO-US042406.

XX

PR 22-DEC-2003; 2003US-0531809P.

XX

PA (AVAL-) AVALON PHARM INC.

XX

PI Weigle B, Ebner R;

XX

DR WPI; 2005-497793/50.

<http://es.ScoreAccessWeb/GetItem.action?AppId=10552...0-552-515-1> copy 157 933.rag&ItemType=4&startByte=0 (13 of 42)10/10/2008 8:50:59 AM

Qy	61	LEVVPDVPPEYYSCFRVFNKLPRFLGSDNQDTFFTSTKRHQILFEILAKTPYGHEKKNLL	120
Db	218	LEVVPDVPPEYYSCFRVFNKLPRFLGSDNQDTFFTSTKRHQILFEILAKTPYGHEKKNLL	277
Qy	121	GIHQLLAEGVLSAAFPLHDGPFKTPPEGPQAPRLNQQRQVLFQHWARWGKWNKYQPLDHVR	180
Db	278	GIHQLLAEGVLSAAFPLHDGPFKTPPEGPQAPRLNQQRQVLFQHWARWGKWNKYQPLDHVR	337
Qy	181	RYFGEKVVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	240
Db	338	RYFGEKVVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	397
Qy	241	CLDCPFWLLSSACALAQAGRLFHDGGTVFFSLFMALWAVLLLEYWKRKSATLAYRWDCSD	300
Db	398	CLDCPFWLLSSACALAQAGRLFHDGGTVFFSLFMALWAVLLLEYWKRKSATLAYRWDCSD	457
Qy	301	YEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVMCLV	360
Db	458	YEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVMCLV	517
Qy	361	SIILYRAIMAIVVSRSGNTLLAAWASRIASLTGSGVNVLFILILSKIYVSLAHVLRWEM	420
Db	518	SIILYRAIMAIVVSRSGNTLLAAWASRIASLTGSGVNVLFILILSKIYVSLAHVLRWEM	577
Qy	421	HRTQTKFEDAFTLKVFIFQFVNFISSPVYIAFFKGRFVGYPGNYHTLFGVRNEECAAGGC	480
Db	578	HRTQTKFEDAFTLKVFIFQFVNFISSPVYIAFFKGRFVGYPGNYHTLFGVRNEECAAGGC	637
Qy	481	LIELAQELLVIMVGKQVINNMQEVLIPLKLGWWQKFLRLSKKRKAGASAGASQGPWEDDY	540
Db	638	LIELAQELLVIMVGKQVINNMQEVLIPLKLGWWQKFLRLSKKRKAGASAGASQGPWEDDY	697
Qy	541	ELVPCEGLFDEYLEMVLQFGFVTIFVAACPLAPLALLNNWVEIRLDARKFVCEYRRPVA	600
Db	698	ELVPCEGLFDEYLEMVLQFGFVTIFVAACPLAPLALLNNWVEIRLDARKFVCEYRRPVA	757
Qy	601	ERAQDIGIWFHILAGLTHLAVISNAFLAFSSDFLPRAYYRWTRAHDLRGFLNFTLARAP	660
Db	758	ERAQDIGIWFHILAGLTHLAVISNAFLAFSSDFLPRAYYRWTRAHDLRGFLNFTLARAP	817
Qy	661	SSFAAAHNRTC	671
Db	818	SSFAAAHNRTC	828

RESULT 5

ABG15488

ID ABG15488 standard; protein; 898 AA.

XX

AC ABG15488;  
 XX  
 DT 18-FEB-2002 (first entry)  
 XX  
 DE Novel human diagnostic protein #15479.  
 XX  
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;  
 KW food supplement; medical imaging; diagnostic; genetic disorder.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200175067-A2.  
 XX  
 PD 11-OCT-2001.  
 XX  
 PF 30-MAR-2001; 2001WO-US008631.  
 XX  
 PR 31-MAR-2000; 2000US-00540217.  
 PR 23-AUG-2000; 2000US-00649167.  
 XX  
 PA (HYSE-) HYSEQ INC.  
 XX  
 PI Drmanac RT, Liu C, Tang YT;  
 XX  
 DR WPI; 2001-639362/73.  
 DR N-PSDB; AAS79675.  
 XX  
 PT New isolated polynucleotide and encoded polypeptides, useful in  
 PT diagnostics, forensics, gene mapping, identification of mutations  
 PT responsible for genetic disorders or other traits and to assess  
 PT biodiversity.  
 XX  
 PS Claim 20; SEQ ID NO 45847; 103pp; English.  
 XX  
 CC The invention relates to isolated polynucleotide (I) and polypeptide (II)  
 CC sequences. (I) is useful as hybridisation probes, polymerase chain  
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,  
 CC and in recombinant production of (II). The polynucleotides are also used  
 CC in diagnostics as expressed sequence tags for identifying expressed  
 CC genes. (I) is useful in gene therapy techniques to restore normal  
 CC activity of (II) or to treat disease states involving (II). (II) is  
 CC useful for generating antibodies against it, detecting or quantitating a  
 CC polypeptide in tissue, as molecular weight markers and as a food  
 CC supplement. (II) and its binding partners are useful in medical imaging  
 CC of sites expressing (II). (I) and (II) are useful for treating disorders  
 CC involving aberrant protein expression or biological activity. The  
 CC polypeptide and polynucleotide sequences have applications in  
 CC diagnostics, forensics, gene mapping, identification of mutations  
 CC responsible for genetic disorders or other traits to assess biodiversity

CC and to produce other types of data and products dependent on DNA and  
 CC amino acid sequences. ABG00010-ABG30377 represent novel human diagnostic  
 CC amino acid sequences of the invention. Note: The sequence data for this  
 CC patent did not appear in the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences

XX

SQ Sequence 898 AA;

Query Match 73.5%; Score 3031.5; DB 4; Length 898;  
 Best Local Similarity 91.2%; Pred. No. 6.3e-316;  
 Matches 578; Conservative 2; Mismatches 11; Indels 43; Gaps 2;

Qy	1	QQDVQDGNNTTVHYALLSASWAVLCYYAEDLRRLKPLQELPNQASNWSAGLLAWLGIPNVL	60
Db	250	QQDVQDGNNTTVHYALLSASWAVLCYYAEDLRRLKPLQDYPTRPPTGRPACCAWLGPINVL	309
Qy	61	LEVVPDVPPEYYSCRFRVKNLPRFLGSDNQDTFFTSTKRHQILFEILAKTPYGHEKKNLL	120
Db	310	LEVVPDVPPEYYSCRFRVKNLPRFLGSDNQDTFFTSTKRHQILFEILAKTPYGHEKKNLL	369
Qy	121	GIHQLLAEGVLSAAFLPHDGFPTPEEGPQAPRLNQRQVLFQHWARWGKWNKYQPLDHVR	180
Db	370	GIHQLLAEGVLSAAFLPHDGFPTPEEGPQAPRLNQRQVLFQHWARWGKWNKYQPLDHVR	429
Qy	181	RYFGEKVALYFAWLGFYTGWLLPAAVVGTIVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	240
Db	430	RYFGEKVALYFAWLGFYTGWLLPAAVVGTIVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	489
Qy	241	CLDCPFWLLSSACALAQ----AGRLFDHGGTVFFSLFMALWAVLLLEYWKRKSATLAYRW	296
Db	490	CLDCPFWLLSSACALAQVREEAGRLFDHGGTVFFSLFMALWAVLLLEYWKRKSATLAYRW	549
Qy	297	DCSDYEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVV	356
Db	550	DCSDYEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMVAVVV	609
Qy	357	MCLVSIILYRAIMAIVSRSGNTLLAAWASRIASLTGSVVNLVFIILSKIYVSLAHVLT	416
Db	610	MCLVSIILYRAIMAIVSRSGNTLLAAWASRIASLTGSVVNLVFIILSKIYVSLAHVLT	669
Qy	417	RWEMHRTQTKFEDAFTLKVFIFQFVNFYSSPVYIAFFKGRFVGYPGNYHTLFGVRNEECA	476
Db	670	RWEMHRTQTKFEDAFTLKVFIFQFVNFYSSPVYIAFFKGRFVGYPGNYHTLFGVRNEECA	729
Qy	477	AGGCLIELAQELLVIMVGKQVINNMQEVFLIPKLKGWQKFRRLRSKKRKAGASAGASQGPW	536
Db	730	AGGCLIELAQELLVIMVGKQVINNMQEVFLIPKLKGWQKFRRLRSKKRKAGASAGASQGPW	789



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DR PC:NCBI; gi30520318.

XX

PT New polynucleotides and polypeptides, useful for developing a diagnostic  
PT marker or medicines for regulation of their expression and activity, or  
PT as targets of gene therapy.

XX

PS Claim 1; Page; 222pp; English.

XX

CC The invention discloses a polynucleotide comprising a sequence selected  
CC from 1970 fully defined nucleotide sequences which encode novel  
CC polypeptides. Also claimed is a polypeptide encoded by the polynucleotide  
CC or its partial peptide, an antibody binding to the polypeptide or peptide  
CC of the polynucleotide, immunologically assaying the polypeptide or  
CC peptide of the polynucleotide by contacting the polypeptide or peptide  
CC with the antibody of the encoded protein, and observing the binding  
CC between the two, a transformant carrying the polynucleotide in an  
CC expressible manner and an antisense polynucleotide. The oligonucleotide  
CC is useful as a primer for synthesising the polynucleotide, or as a probe  
CC for detecting the polynucleotide. The polynucleotides and encoded  
CC proteins are useful as pharmaceutical agents and many disease-related  
CC genes may be included in them, for developing a diagnostic marker or  
CC medicines for regulation of their expression and activity, or as targets  
CC of gene therapy. The genes are involved in tissue and/or cell  
CC regeneration. Membrane proteins, signal transduction-related proteins,  
CC transcription-related proteins, disease-related proteins and genes  
CC encoding them can be used as indicators for diseases (e.g. osteoporosis,  
CC neurological diseases, cancer, tumours. The cDNA may be used to regulate  
CC the activity or expression of the encoded protein to treat diseases. The  
CC sequence presented is a protein of the invention. Note: Some of the  
CC sequence data for this patent is not represented in the printed  
CC specification, but is based on sequence information supplied by the  
CC European Patent Office.

CC

CC Revised record issued on 15-JUN-2007 : Enhanced with precomputed  
CC information from BOND.

XX

SQ Sequence 920 AA;

Query Match 36.4%; Score 1502.5; DB 6; Length 920;  
Best Local Similarity 40.4%; Pred. No. 2.5e-151;  
Matches 328; Conservative 145; Mismatches 270; Indels 69; Gaps 20;

Qy 8 NTTVHYALLSASWAFLCYAEDLRLKLPQE---LPNQASNWS-----AGLLAWLGIP 57  
| : : : | | | | | : : : : : | : | : | |  
Db 122 NSDIIFVKLHAPWEVLGRYAEQMNVRMPFRRKIYYLPRRYKFMSRIDKQISRLRRWLPKK 181  
  
Qy 58 NVLL--EVVPDVPP-EYYSRFRVKNLPRFLGSDNQDTFTTSTKRHQILFEILAKTPYGH 114  
: | | : | : : : : : | : : : | : : : | : : : |  
Db 182 PMRLDKETLPDLEENDCYTAPFSQQRHHFI-IHNKETFFNNATRSRIVHHILQRIKY-E 239

Qy	115	EKKNLLGIHQLLAEGVLSAAFLPHDGPFFKTPPEGPQAPRLNQRVLFQHWARWGKWNKYQ	174
Db	240	EGKNKIGLNRLLTNGSYEAAFLHEGSYRSKNSIRTHGAENHRHLLYECWASWGVVYKYQ	299
Qy	175	PLDHVRRYFGEKVALYFAWLGFYTGWLLPAAVVGTLVLVLVCGFLVFSDIPTQELCGSKDS	234
Db	300	PLDLVRRYFGEKIGLYFAWLGWYTGMFLPAAFIGLFVFLYGVTTLDHSQVSKEVCQATDI	359
Qy	235	FEMCPLC-LDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAVLLEWYWRKSATLA	293
Db	360	I-MCPVCDKYCPFMRLSDSCVYAKVTHLFDNGATVFFAVFMAVWATVLEFWKRRRAVIA	418
Qy	294	YRWDCSDYEDTEERPRPQFAAS-APMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMV	352
Db	419	YDWDLIDWEEEEERIPQFEAKYSKKERNMNPISGKPEPYQAFTDKCSRLIVSASGIFFMI	478
Qy	353	AVVVMCLVSIILYRAIMAIIVSRSGNTLLA-AWA-----SRIASLTGSGV--NLVFILIL	404
Db	479	CVVIAAVFGIVIRYRVTV-----STFAAFKVALIRNNSQVAT-TGTAVCNFCIIMLL	530
Qy	405	SKIYVSLAHVLTRWEMHRTQTKFEDAFTLKVFIQFVNFISSPVYIAFFKGRFVGYPGNY	464
Db	531	NVLYEKVALLLTNLEQPRTESEWENSFTLKMFLQFQVNLNSSTFYIAFFLGRFTGHGPAY	590
Qy	465	HTLFG-VRNEECAAGGLIELAQELLVIMVGKQVINNMQEVLIPLKLGWWQKFLRLSKKR	523
Db	591	LRLINRWRLEECHPSGCLIDLQMGIMVLKQTNWNNFMELGYPLIQNWTR---RKVRQ	647
Qy	524	KAGASAGASQGPWEDDYELVPC--GLFDEYLEMVLQGFVTFVAAACPLAPLALLNNW	581
Db	648	EHGPERKISFPQWEKDYNLQPMNAYGLFDEYLEMILQGFTTTFVAAFPPLAPLLALLNNI	707
Qy	582	VEIRLDARKFVCEYRRPVAERAQDIGIWFHILAGLTHLAVISNAFLLAFSSDFLPRAYR	641
Db	708	IEIRLDAYKFVTQWRRLASRAKDIGIWIYGILEGIGILSVITNAFVIAITSDFIPLVYA	767
Qy	642	W-----TRAHDLRGFLNFTLA-----RAPSSFAAAHNRTCRYAFR	677
Db	768	YKYGPCAGQGEAGQKCMVGYVNASLSVFRISDFENRSEPSDGESEFSGTPLYCRYRDR	827
Qy	678	DDDGH----YSQTYWNLLAIRLAFVIVFEHVVSFVGRLLDLLVPDIPESVEIKVKREYY	732
Db	828	DPPHSLVPYGYTLQFVHWLAARLAFIIVFEHLVFCIKHLISYLIPLDKDLRDRMRREKY	887
Qy	733	LAKQALAENEVLFGTNGTKDEQPKGSELSSHW	764
Db	888	LIQEMMYEAELERLQKERRKKNKKAHHNEW	919

RESULT 7

ABP58666

ID ABP58666 standard; protein; 920 AA.

XX

AC ABP58666;

XX

DT 24-MAR-2003 (first entry)

XX

DE Human dihydropyrimidinase related protein 1-101.20.

XX

KW Human; dihydropyrimidinase related protein 1-101.20;

KW recombinant production; gene therapy; psychosis; development disorder;

KW uracil-related metabolic disorder; thymine-related metabolic disorder;

KW pyrimidine metabolic disorder.

XX

OS Homo sapiens.

XX

PN CN1364894-A.

XX

PD 21-AUG-2002.

XX

PF 10-JAN-2001; 2001CN-00105195.

XX

PR 10-JAN-2001; 2001CN-00105195.

XX

PA (BIOW-) BIOWINDOW GENE DEV INC SHANGHAI.

XX

PI Mao Y, Xie Y;

XX

DR WPI; 2003-000532/01.

DR N-PSDB; ABZ57080.

XX

PT New polypeptide-human dihydropyrimidinase relative protein 1-101, 20 and  
PT polynucleotide for encoding such polypeptide.

XX

PS Claim 1; Page 28-30 (Disclosure); 36pp; Chinese.

XX

CC The invention relates to human dihydropyrimidinase related protein 1-  
CC 101.20 (ABP58666) and nucleic acids encoding it (ABZ57080). The protein  
CC has a molecular weight of 101.2 kD. The invention also relates to a  
CC method for the recombinant production of the protein, an antagonist of  
CC the protein, and the use of the protein, gene and antagonist in  
CC therapeutic applications. Dihydropyrimidinase related protein 1-101.20  
CC can be used in the treatment of a variety of diseases such as psychosis,  
CC development disorders and uracil- and thymine-related metabolic  
CC disorders. The present sequence represents human dihydropyrimidinase  
CC related protein 1-101.20

XX

SQ Sequence 920 AA;

Query Match 36.0%; Score 1482.5; DB 6; Length 920;  
 Best Local Similarity 40.0%; Pred. No. 3.6e-149;  
 Matches 325; Conservative 146; Mismatches 272; Indels 69; Gaps 20;

Qy	8	NTTVHYALLSASWAFLCYAEDLRKLPLQE----	LPNQASNWS-----AGLLAWLGIP	57
		: : :               : : :   :   :		
Db	122	NSDIIFVKLHAPWEVLGRYAEQMNVRMPFRRKIYYLPRRYKFMSRIDKQISRFRRLWPKK		181
Qy	58	NVLL--EVVPDVP--EYYSRFRVFNKLPRFLGSDNQDTFTSTKRHQLFEILAKTPYGH		114
		:     :   : : :   :   : :   :   :   :   :   :		
Db	182	PMRLDKETLPDLEENDCYTAPFSQQRIHHFI-IHNKETTFNNATRSRIVHHILQRIKY-E		239
Qy	115	EKKNLLGIHQLLAEGVLSAAFLPHDGPFFKTPPEGPQAPRLNQQRVLFQHWARWGKWNKYQ		174
		: : :             : : :     : : :		
Db	240	EGKNKIGLNRLLTNGSYEAAFLHEGSYRSKNSIRTHGAENHRHLLYECWASWGVWYKYQ		299
Qy	175	PLDHVRRYFGEKVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDS		234
		:         :       :       : : :   :		
Db	300	PLDLVRRYFGEKIGLYFAWLGWYTGMFLPAAFIGLFVFLYGVTTLDHSQVSKEVCQATDI		359
Qy	235	FEMCPLC-LDCPFWLLSSACALAQAGRLFDHGGTVFFSFLMALWAVLLLEYWKRKSATLA		293
		:         :   :     :     :     :   :     :		
Db	360	I-MCPVCDKYPFMRSLSDSCVYAKVTHLFDNGATVFFAFVMAVWATVFLFWKRRRAVIA		418
Qy	294	YRWDCSDYEDTEERPRPQFAAS-APMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMV		352
		: :         :     :     :   : :   :		
Db	419	YDWDLIDWEEEEEEIRPQFEAKYSKKERNMNPISGKPEPYQAFTDKCSRLIVSASGIFFMI		478
Qy	353	AVVVMCLVSIILYRAIMAIVVSRSGNTLLA-AWA-----SRIASLTGSVV--NLVFILIL		404
		: :   :   : :         : :     :     :		
Db	479	CVVIAAVFGIVIRVTV-----STFAAFKVALIRNNSQVAT-TGTAVCINFCIIMLL		530
Qy	405	SKIYVSLAHVLTRWEMHRTQTKFEDAFTLKVFIQFVNFIYSSPVYIAFFKGRFVGYPGNY		464
		: :   :   :       : : : :     :		
Db	531	NVLYEKVALLLTNLEQPRTESEWENSFTLKMFLFQFVNLSSTFYIAFFLGRFTGHGPAY		590
Qy	465	HTLFG-VRNEECAAGGLIELAQELLVIMVGKQVINNMQEVFLIPKLKGWVQKFLRLSKKR		523
		: : :           : : :   : :   : :		
Db	591	LRLINRWRLLEECHPSGLIDLQCMQIMVLKQTWNNFMELGYPLIQNWVTR---RKVRQ		647
Qy	524	KAGASAGASQGPWEDDYELVPC--GLFDEYLEMVLQFGFVTIFVAACPLAPLALLNNW		581
		:             :       :		
Db	648	EHGPERKISFPQWEKDYNLQPMNAYGLYDEYLEMILQFGFTTIFVAAFPPLAPLLALLNNI		707
Qy	582	VEIRLDARKFVCEYRRPVAERAQDIGIWFHILAGLTHLAVISNAFLAFSSDFLPRAYYR		641
		:               : :   :     :     :   :   :   :		
Db	708	IEIRLDAYKFVTQWRRPLASRAKDIGIGYGILEGIGILSVITNAFVIAITSDFIPRLVYA		767

Qy 642 W-----TRAHDLRGFLNFTLA-----RAPSSFAAAHNRTCRYRAFR 677  
 : : |:| :|: | |: : ||| :|  
 Db 768 YKYGPCAGQGEAGQKCMVGYVNASLSVFRISDFENRSEPESDGSEFSGTPLYCRYRDYR 827

Qy 678 DDDGH----YSQTYWNLLAIRLAFVIVFEHVVSFVGRLLDLLVPDIPESVEIKVKREYY 732  
 | : :|:| | | | :| | : | :| :| : :| :|  
 Db 828 DPPHSLVPYGYTLQFVHVLARLAFIIVFEHLVFCIKHLISYLIPDLPKDLRDMRREKY 887

Qy 733 LAKQALAENEVLFGTNGTKDEQPKGSELSSHW 764  
 | : : | | : | : | : |  
 Db 888 LIQEMMYEAELERLQKERKERKNGKAHHNEW 919

## RESULT 8

ADK52114

ID ADK52114 standard; protein; 981 AA.

XX

AC ADK52114;

XX

DT 15-JUN-2007 (revised)

DT 20-MAY-2004 (first entry)

XX

DE Human atopic dermatitis/psoriasis-associated protein #29.

XX

KW Human; atopic dermatitis; psoriasis; dermatological; anti-inflammatory;

KW antipsoriatic; rash; BOND\_PC; transmembrane protein 16C;

KW chromosome 11 open reading frame 25;

KW transmembrane protein 16C [Homo sapiens]; TMEM16C; C11orf25; GENX-3947;

KW transmembrane protein 16C (eight membrane-spanning domains);

KW hypothetical protein; hypothetical protein [Homo sapiens]; GO16020;

KW GO16021; GO4185; GO7001.

XX

OS Homo sapiens.

XX

PN WO2004016785-A1.

XX

PD 26-FEB-2004.

XX

PF 06-AUG-2003; 2003WO-JP009999.

XX

PR 06-AUG-2002; 2002JP-00229319.

PR 14-MAY-2003; 2003JP-00136544.

XX

PA (GENO-) GENOX RES INC.

PA (UYJU-) UNIV JUNTENDO.

XX

PI Itoh M, Ogawa K, Shinagawa A, Sudo H, Ogawa H, Ra C;

PI Mitsuishi K;

PT Detecting atopic dermatitis or psoriasis comprises assaying levels of  
PT expression of an indicator gene at a rash site and non-rash site of a  
PT person with atopic dermatitis or psoriasis.

CC The invention relates to detecting atopic dermatitis or psoriasis  
CC comprising assaying the levels of expression of an indicator gene at a  
CC rash site and non-rash site of a person with atopic dermatitis or  
CC psoriasis, comparing these levels with those of a healthy person, and  
CC determining that if the levels of indicators are higher or lower, then  
CC this indicates the disease. Also included are a reagent for detecting  
CC atopic dermatitis or psoriasis, a kit for screening for treatments, a  
CC transgenic non human vertebrate animal models for the diseases, an agent  
CC for inducing the diseases in mice and a DNA chip for assaying for the  
CC indicator genes. The method is used for treatment, detection and animal  
CC models for research of atopic dermatitis and psoriasis. The present  
CC sequence is a protein encoded by an indicator gene of the invention.

SO Sequence 981 AA;

Query Match 35.6%; Score 1467.5; DB 8; Length 981;  
Best Local Similarity 40.9%; Pred. No. 1.6e-147;  
Matches 313; Conservative 149; Mismatches 245; Indels 59; Gaps 21;

Qy 20 WAVLCYYAEDRLRLKLPLQ-----ELPNQASNWSAGLLAWLGIPNVLLE--VVPDV 67  
| | | | | : : : : : : : : : : : : : : : : : :  
Db 214 WDTLCKYAEARNIRMFFRKCKYYTDGRSKSMGRMOTYFRRIKDWMAONPMVLDKSAFPDL 273

Qy           68 --PPEYYSRFRVNKLPRFLGSDNQDTFFFTSTKRHQILFEILAKTPY--GHEKKNLLGIHQ 124  
       | : | : | : | : | : | : | : | : | : | : | : | : | : | : | : | :  
Db          274 EESDCYTGPFSRARIRHHFI-INNKDTFFSNATRSRVVHMLERTKYENGISK---VGIR 329

Qy 125 LLAEGVLSAAFLPHDGFPKT----PPEGPAQAPRLNQRQVLFGQHWAARWGKWNKYQLPDHVR 180  
| : | : | : | : | : | : | : | : | : | : | : | : | : | : | : |  
Db 330 LINNGSYIAAFPPHEGAYKSSOPIKTHGP-----NNRHLLYERWARGWMWYKHOPDLIR 385

Qy 181 RYFGERVALYFAWLGFYTGWLLPAAVVGLTVLVLGVCFLVFSDIPTQELCGSKDSFEMCPL 240  
 |||||:|||||:|||||:|||||:| | | : : : :||| : : | |||

Db 386 LYFGEGIGLYFAWLGWYTGMLIPAAIVGLCVFFYGLFTMNNSOVSQICKAEVFCMPL 444

<http://es.ScoreAccessWeb/GetItem.action?AppId=10552...0-552-515-1> copy 157 933.rag&ItemType=4&startByte=0 (24 of 42)10/10/2008 8:50:59 AM



KW diagnostic; metastasis; esophagus tumor; gastrointestinal disease;  
 KW neoplasm; cytostatic; cancer; AXL; ZBTB11; TNFRSF14; NSUN5; SPEN; LTBP3;  
 KW SYNGR1; SLC13A1; MAP3K12; GLYAT; ZNF659; B4GALT2; POGK; AQP3; CAPG;  
 KW SLIT2; BOND\_PC; transmembrane protein 16C;  
 KW chromosome 11 open reading frame 25;  
 KW transmembrane protein 16C [Homo sapiens]; TMEM16C; C11orf25; GENX-3947;  
 KW transmembrane protein 16C (eight membrane-spanning domains);  
 KW hypothetical protein; hypothetical protein [Homo sapiens]; GO16020;  
 KW GO16021; GO4185; GO7001.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO2006118308-A1.  
 XX  
 PD 09-NOV-2006.  
 XX  
 PF 02-MAY-2006; 2006WO-JP309177.  
 XX  
 PR 02-MAY-2005; 2005JP-00134530.  
 PR 13-SEP-2005; 2005JP-00265645.  
 PR 13-SEP-2005; 2005JP-00265678.  
 XX  
 PA (TORA ) TORAY IND INC.  
 PA (KYOU ) UNIV KYOTO.  
 XX  
 PI Akiyama H, Kozono S, Myomoto A, Nomura O, Nobumasa H, Tanaka Y;  
 PI Tomoda S, Shimada Y, Tsujimoto G;  
 XX  
 DR WPI; 2007-110304/11.  
 DR PC:NCBI; gil13899227.  
 DR PC:SWISSPROT; Q9BYT9.  
 XX  
 PT Composition for determining occurrence/metastasis of esophageal cancer in  
 PT subject, comprises an antibody binding to a polypeptide encoded by a  
 PT polynucleotide having a sequence of genes e.g. AXL, ZBTB11 and TNFRSF14,  
 PT and/or polynucleotides.  
 XX  
 PS Claim 1; SEQ ID NO 231; 142pp; Japanese.  
 XX  
 CC This invention describes a novel composition for detecting metastasis of  
 CC esophageal cancer in a test subject. The composition contains a probe  
 CC derived from polynucleotides AXL, ZBTB11, TNFRSF14, NSUN5, SPEN, LTBP3,  
 CC SYNGR1, SLC13A1, MAP3K12, GLYAT, ZNF659, B4GALT2, POGK, AQP3, CAPG,  
 CC SLIT2, their variants or fragments and an antibody. The invention also  
 CC claims: a) a kit for detecting, determining or presuming the occurrence  
 CC or metastasis of esophageal cancer in a test subject; b) a DNA chip for  
 CC detecting, determining or presuming the occurrence or metastasis of  
 CC esophageal cancer and c) a method to detect, determine or presume the  
 CC occurrence or metastasis of esophageal cancer in a test subject by

CC detecting the presence of or amount or expression level of one or more  
 CC esophagus-cancer related target nucleic acid in a biological sample. The  
 CC method enables the rapid and convenient detection of occurrence or  
 CC metastasis of esophageal cancer in test subject with high sensitivity.  
 CC This sequence represents a protein used in the method of the invention  
 CC  
 CC Revised record issued on 15-JUN-2007 : Enhanced with precomputed  
 CC information from BOND.

XX

SQ Sequence 981 AA;

Query Match 35.6%; Score 1467.5; DB 12; Length 981;  
 Best Local Similarity 40.9%; Pred. No. 1.6e-147;  
 Matches 313; Conservative 149; Mismatches 245; Indels 59; Gaps 21;

Qy	20	WAVLCYYAEDLRKLPLQ-----ELPNQASNWSAGLLAWLGIPNVLE--VVPDV	67
		: : : : :   : :   :	
Db	214	WDTLCKYAEARNLIRMPFRKKCCYYTDGRSKSMGRMQTYFRRIKDWMAQNPMLDKSAFPDL	273
Qy	68	-PPEYYSCRFRVNKLPRFLGSDNQDTFTTSTKRHQILFEILAKTPY--GHEKKNLLGIHQ	124
		:   :   : :   : :   :   :   :   :   :   :	
Db	274	EESDCYTGPFSSRARIHFI-INNKDTFFSNATRSRIVYHMLERTKYENGISK---VGIRK	329
Qy	125	LLAEGVLSAAFPPLHDGPFT----PPEGPQAPRLNQRQVLFQHWARWGKWNKYQPLDHVR	180
		:         :   : :         : :   :   :   :   :	
Db	330	LINNGSYIAAFPPEGAYKSSQPIKTHGPQ---NNRHLLYERWARWGMWYKHQPLDLIR	385
Qy	181	RYFGEKVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKDSFEMCPL	240
		:         :     :         :   :   :   :	
Db	386	LYFGEKIGLYFAWLGWYTGMLIPAAIVGLCVFFYGLFTMNNSQVQSQICKATEVF-MCPL	444
Qy	241	C-LDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAVLLLEYWKKRSATLAYRWDCS	299
		:   : :   :   :     :     :     :   :   :   :	
Db	445	CDKNCSLQRLNDSCIYAKVTYLFDNGGTVFFAIFMAIWAIVLEFWKRRRSILTYTWDLI	504
Qy	300	DYEDTEERPRPQFAAS-APMTAPNPITGEDEPYFPERSARRMLAGSVVIVVMVAVVMC	358
		: :   :             :         :   :   :   :   :   :	
Db	505	EWEETLRPQFEAKYYKMEIVNPITGKPEHPQSSDKVTRLVSVSGIFFMISLVITA	564
Qy	359	LVSIIYR-AIMAIVVSRSGNTLLAAWASRIASLTGSV-VNLVFILILSKIYVSLAHVLT	416
		: : :     :   :   :   :   :   :   :   :   :   :	
Db	565	VFGVVVYRLVVMQFASFQWNFQKQY--QFATSAAAVCINFIIIMLLNLAYEKIAYLLT	622
Qy	417	RWEMHRTQTKFEDAFTLKVFIQFVNIFYSSPVYIAFFKGRFVGYPGNHYHTLFG-VRNEEC	475
		: : : : :   :   :                     :	
Db	623	NLEYPRTESEWENSFALKMFLQFVNLNSSIFYIAFFLGRFVGHPGKYNKLFDRWRLEEC	682
Qy	476	AAGGLIELAQELLVIMVGKQVINNMQEVLPKLKGWWQKFLRLSKRRKAGASAGASQGP	535
		:   : :       :     :   :   :   : :	

AFG11146

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Qy	232	KDSFEMCPLC-LDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAVLLLEYWKRKSA	290
		: :         :  :        :      :      :      : :	
Db	257	RHNITMCPLCDKTCSYWKMSACATARASHLFDNPATVFFSVFMALWAATFMEHWKRKQM	316
Qy	291	TLAYRWDCSDYEDTEERPRPQFAA-----SAPMTAPNPITGEDEPYFPERSARRMLAGS	345
		: :  :  :  :  :  :  :  :  :  :  :  :	
Db	317	RLNYRWDLTGFEEDDHPRAEYEARVLEKSLKKESRNKET--DKVKLTWRDRFPAYLTNL	374
Qy	346	VVIVVMVAVVVMCLVSIILYRAIMAIIVVSRSGNTLLAAWASRIASLTGSVVNLVFILILS	405
		:  :	
Db	375	VSIIFMIAVTFaIVLGVIIYRISMAAALAMNSSPSVRSNIRVTVTATAVINLVVILLDD	434
Qy	406	KIYVSLAHVLTRWEMHRTQTKEFADFTLKVFIFQFVNFISSPVYIAFFKGRFVGYPGNYH	465
		:  :	
Db	435	EVYGCiARWLTkIEVPKTEKSFEERLIFKAFLLKFVNSYTPIFYVAFFKGRFVGRPGDYV	494
Qy	466	TLF-GVRNEECAAGGCLIELAQELLVIMVGKQVI--NNMQEVLIPKLGWVQKFRLRSKKR	523
		:  :	
Db	495	YIFRSFRMEECAPGGCLMELCIQLSIIMLGKQLIQNNLFBIGIPKMKKLIRYLKLKQQSP	554
Qy	524	KAGASAGASQGPWEDDYELVPCGLFDEYLEMVLQFGFVTIFVAACPLAPLFPALLNNWVE	583
		: :  :	
Db	555	PDHEECVKKRQRYEVDYNLEPFAGLTPEYMEMIIQFGFVTLFVASFPLAPLFPALLNNIIE	614
Qy	584	IRLDARKFVCEYRRPVAERAQDIGIWFHILAGLTHLAVISNAFLAFSSDFLPRA--YYR	641
		:              :     :    :  :  :  :  :  :  :  :  :  :	
Db	615	IRLDAKKFVTELRRPVAVRAKDIGIWNILRGIGKLAVIIDAFVISFTSDFIPRLVLYLM	674
Qy	642	WTRAHDLRGLNFTLARAPSSF-----AAAHN-----RTCryAFRD---DDGH	682
		: :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :	
Db	675	YSKNGTMHGFVNHTL----SSFNVSDFQNGTAPNDPLDLGYEVQICRYKDYREPPWSENK	730
Qy	683	Y--SQTYWNLLAIRLAFVIVFEHVVSvGRLLDLLVPDIPESVEIKVKREYYLA-----	734
		:  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :	
Db	731	YDISKDFWAVLAARLAFVIVFQNLVFMMSDFVDWVDPIDPKDISQQIHKEKVLMMVLFMR	790
Qy	735	----KQALAENEVLFGTNGTKDEQP-----KGSELSSH	763
Db	791	EEQDKQQLL--ETCMEKERQKDEPPCNHNTKACPDLSGPAPSH	833

# RESULT 11

AEG11142

ID AEG11142 standard; protein; 960 AA.

XX

AC AEG11142;

XX

DT 15-JUN-2007 (revised)

DT 20-APR-2006 (first entry)

XX  
DE Human transmembrane protein 16A, SEQ ID NO: 7.  
XX  
KW Genetic marker; diagnostic; prognosis; gastrointestinal tumor;  
KW cytostatic; neoplasm; tumor marker; transmembrane protein 16A; BOND\_PC;  
KW transmembrane protein 16A;  
KW transmembrane protein 16A (eight membrane-spanning domains);  
KW oral cancer overexpressed 2; membrane protein;  
KW tumor amplified and overexpressed sequence 2;  
KW transmembrane protein 16A [Homo sapiens]; TMEM16A; TAOS2; ORAOV2;  
KW FLJ10261.  
XX  
OS Homo sapiens.  
XX  
PN US2006040292-A1.  
XX  
PD 23-FEB-2006.  
XX  
PF 08-JUL-2005; 2005US-00177894.  
XX  
PR 08-JUL-2004; 2004US-0586676P.  
XX  
PA (WEST/) WEST R B.  
PA (VRIJ/) VAN DE RIJN M.  
XX  
PI West RB, Van De Rijn M;  
XX  
DR WPI; 2006-182760/19.  
DR N-PSDB; AEG11136.  
DR REFSEQ; NP\_060513.  
DR PC:NCBI; gi40354210.  
XX  
PT Classifying tumor as gastrointestinal stromal tumor belonging to PDGFRA  
PT positive subclass, involves detecting expression or activity of gene  
PT encoding DOG1 polypeptide in sample.  
XX  
PS Disclosure; SEQ ID NO 7; 177pp; English.  
XX  
CC The present invention relates to three gene markers such as DOG1, KIT and  
CC platelet derived-growth factor receptor alpha (PDGFRA) that are useful in  
CC classifying tumors. These gene markers are useful in the classification  
CC of gastrointestinal stromal tumors (GISTs) and tumors other than GISTs.  
CC The invention also relates to methods providing diagnostic, prognostic  
CC and predicative information based on the classifying step. The invention  
CC is useful for classifying gastrointestinal stromal tumors as belonging to  
CC a PDGFRA positive subclass, KIT negative or PDGFRA negative subclass. The  
CC present sequence is human transmembrane protein 16A (DOG1; TMEM16A). The  
CC DOG1 gene encodes a transmembrane protein of unknown function  
CC (transmembrane protein 16A). The transmembrane protein 16A is encoded by



Db 671 QQSPPDHEECVKRKQRYEVDYNLEPFAGLTPEYMEMI IQFGFVTLFVASFLAPLFLALLN 730

Qy 580 NWVEIRLDARKFVCEYRRPVAERAQDIGIWPHILAGLTHLAVISNAFLAFSSDFLPRA- 638  
| :|||||:| | | |||| ||:|||||:| | | : |||| |||:|:|:|:| |

Db 731 NIIIEIRLDAKKFVTELRRPVAVRADIGIWIYNILRGIGKLAVIINAFVISFTSDFIPRLV 790

Qy 639 -YYRWTRAHDLRGFLNFTLARAPSSF-----AAAHN-----RTCryRAFRD--- 678  
| : : : | : | : | | | | | | | : | | : | | : | : :

Db 791 YLYMYSKNGTMHGFVNHTL----SSFNVSDFQNGTAPNDPLDLGYEVQICRYKDYREPPW 846

Qy 679 DDGHY--SQTYWNLLAIRLAFVIVFEHVVSFVGRLDLLVDPDIPESVEIKVKREYYLA-- 734  
: | | : : | : | | | | | | : : : | : : | : : : | : : |

Db 847 SENKYDISKDFWAVLAARLAFVIVFQNLVFMMSDFVDWVIPDIPKDISQQIHKEKVLMEV 906

Qy 735 -----KQALAENEVLFGTNGTKDEQP-----KGSELSSH 763  
| | | | | | | | | | | | | | | | | |

Db 907 LFMREEQDKQQLL--ETWMEKERQKDEPPCNHNTKACPDSLGSPPASH 953

# RESULT 12

AFB77190

ID AFB77190 standard; protein; 1017 AA.

XX

AC AFB77190;

XX

DT 28-JUN-2007 (first entry)

XX

DE Mouse TM-1 (Tmem16a) protein.

XX

KW Cell isolation; stem cell; therapeutic; transgenic animal; screening;  
KW tissue regeneration; genitourinary disease; uropathic;  
KW intervertebral disk displacement; degeneration; injury; vulnerary;  
KW back pain; transmembrane factor-1; Tmem16a.

XX

OS Mus musculus.

XX

PN W02007027583-A2.

XX

PD 08-MAR-2007.

XX

PF 28-AUG-2006; 2006WO-US033491.

XX

PR 31-AUG-2005; 2005US-0713400P.

XX

PA (UYFL ) UNIV FLORIDA RES FOUND INC.

XX

PI Harfe BD, Cohn MJ;

XX

DR WPI; 2007-412931/39.



DR N-PSDB; AFB77189.  
 XX  
 PT Isolating sonic hedgehog expressing-cells comprises obtaining a non-human  
 PT transgenic subject in which a marker gene has been inserted into the  
 PT subject's genome.  
 XX  
 PS Disclosure; SEQ ID NO 2; 96pp; English.  
 XX  
 CC The present invention relates to a method of isolating cells in selected  
 CC tissues co-expressing the sonic hedgehog (Shh) gene and a marker gene.  
 CC The method involves obtaining a non-human transgenic subject in which a  
 CC marker gene has been inserted into the subject's genome and isolating  
 CC Shh/marker gene expressing cells and Shh/marker gene non-expressing cells  
 CC from the selected tissue. The invention further provides a method of  
 CC identifying differentially expressed genes (e.g. transmembrane factors TM  
 CC -1 and TM-2, EST 1437418, Mmu-miR-135a-2 and AP-2 beta) in selected  
 CC tissues co-expressing the sonic hedgehog gene and a marker gene. The  
 CC invention is useful in tissue engineering, regeneration, reconstruction  
 CC and/or repair of tissues and genitourinary system and also in treating  
 CC intervertebral disk rupture, degeneration, disease or injury and back  
 CC pain. The invention is further useful for generating transgenic animal.  
 CC The present sequence is the mouse TM-1 (Tmem16a) protein.  
 XX  
 SQ Sequence 1017 AA;

Query Match 35.2%; Score 1452.5; DB 12; Length 1017;  
 Best Local Similarity 40.0%; Pred. No. 7e-146;  
 Matches 330; Conservative 156; Mismatches 257; Indels 81; Gaps 22;

```

Qy      6  DGNITVH---YALLSASWAVLCYYAEDLRKLKPLQELPNQASNWSAGLLAWLGIPNVLLE 62
      | : | : | : | | | | | | : | : | : : : : : | | | | : | :
Db      202 DEDTKIHGVGVFKIHAPWHVHLCREAEFLKCLKMPTKKVYHISE--TRGLLK--TINSVLQK 257

Qy      63  VVPDVPPEYYSCRFRVNKLPRFLGS-----DNQDIFTTSTKRHQILFEILAKTPYG 113
      : : | : | : | : | : | : | : | : | : | : | : | : |
Db      258 ITDPIQPKVAEHRPQTTRKLSYPFSREKQHLFDLTDSDFFDSKTRSTIVYEILKRTTCT 317

Qy      114 HEKKNLLGIHQLLAEGVLSAAFFPLHDGPFKTPPEGPAQLRNQQLVLFQHWARWGKWNKY 173
      | : : | | | | | | | | | | : | | | : | : | : | : |
Db      318 KAKYS-MGITSLLANGVYSAAYPLHDGDY----EGDNV-BFNDKLLYEAWASYGVFYKY 371

Qy      174 QPLDHVRRYFGEKVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSKD 233
      | | : | | : | | | | | | | | | | : | : | : | : | : | :
Db      372 QPIDLVRKYFGEKVGLYFAWLGAYTQMLIPASIVGVIVFLYGATVDENIPSMEMCDQRY 431

Qy      234 SFEMCPLC-LDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAVLLEYWKRKSATL 292
      : | | | | | | : | : | | | | | : | : | : | : | : | : |
Db      432 NITMCPLCDKTCYSWKMSACATARASHLFDNPAIVFFSVFMALWAATFMEHWKRKQMRL 491

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Qy	168	GKWNKYQPLDHVRRYFGEKVALYFAWLGFTYTWLLPAAVVGTIVFLVGCFLVFSDIPTQE	227
		:  : :  : : : :                :  : : :  : :    :    : :	
Db	341	GVFYKQPIDLIRKYFGEKIGLYFAWLGLYTSFLIPSSVIGVIVFLYGCATIEEDIPSRE	400
Qy	228	LCGSKDSFEMCPLC-LDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAVLLEYYWK	286
		:  :::          :                  :                :	
Db	401	MCDQQNAFTMCPCLCDKSCDYWNLSACGTAQASHLFDNPATVFFSIFMALWATMFLENWK	460
Qy	287	RKSATLAYRWDCSDYEDTEER----PRPQFAA-----SAPMTAPNPIT----G	326
		:   :         :	
Db	461	RLQMRLGYFWDLTGIEEEEEERAQEHRSPEYETKVREKMLKESNQSAVQKLETNTTECGDE	520
Qy	327	EDEPYFPERSRARMLAGSVVIVVMVAVVMCLVSIILYRAIMAIIVSRSGNTLLAAWAS	386
		:    :       : : :  : : :    :	
Db	521	DDEDKLTWKDRFPGYLMNFASILFMIALTFSIVFGVIVYRITTAALS-----LNKATRS	575
Qy	387	RI---ASLTGSVNLVFIILSKIYVSLAHVLTREWHRHTQTKFEDAFTLKVFIQFVNF	443
		: :   : :      :  :  :    :  :  :  :      :  :	
Db	576	NVRVTVTATAVINLVILDEIYGAVAKWLTKIEVPKTEQTFEERLILKAFLLLKFVNA	635
Qy	444	YSSPVYIAFFKGRFVGYPGNYHTLF-GVRNEECAAGGCLIELAQELLVIMVGKQVI--NNM	501
		: : : :      :  :                :  : : : :      :	
Db	636	YSPIFYVAFFKGRFVGPGSVYVYFDGYRMEECAPGGCLMELCIQLSIIMLGKQLIQNNI	695
Qy	502	QEVLPKLGWWQKFLRLRSKKRKAGASAGA-SQGP--WEDDYELVPCEGLFDEYLEMVLQ	558
		: : :  :   : : :  :   :  :   :  :             : : :	
Db	696	FEIGVPKLK---KLFRKLKDETEAGETDSAHSKHPEQWDLDSLEPYTGLTPEYMEMIIQ	752
Qy	559	FGFVTIFVAACPLAPLAFALLNNWVEIRLDARKFVCEYRPRVAERAQDIGIWFHILAGLTH	618
		: :  : : : :  : : : :            : : :      : :	
Db	753	FGFVTLFVASFPLAPVFAALLNNVIEVRLDAKKFVIELRRPDVARTKDIGIWFIDILSGIGK	812
Qy	619	LAVISNAFLAFSSDFLPRAYRWTRAHD--LRGFLNFTLA-----RAPSSFAA	665
		:        : :  : : :    ::: :  :  :  :  :  :  :  :	
Db	813	FSVISNAFVIAITSDFIPRLVQYQSYSHNGTLHGFEVNHLSFFNVSQLKEGTQPENSQFD	872
Qy	666	AHNRTCRYRAFRD-----DDGHYSQTYWNLLAIRLAFVIVFEHVVSFVGRLLDLLVPDIP	720
		:    :  : :  : : :  :  :  :  :              :  :  : :	
Db	873	QEVQFCRFKDYREPPWAPNPYEFKQYWFILSARLAFVIFQNLVMFLSVLVDWMIPDIP	932
Qy	721	ESVEIKVKRE-----YYLAKQALAENEVLFGTNGTKDEQPKGSELSSHWPFTVPKA-S	773
		: : :  : :  :  :  :  :  :  :  :  :  :  :  :  :	
Db	933	TDISDQIKKEKSLLVDFFLKE----EHEKCLKMDEPALRSPGGGDRSRSRAASSAPSGQS	988
Qy	774	QL 775	
Db	989	QL 990	

## RESULT 14

AEH82071

ID AEH82071 standard; protein; 913 AA.

XX

AC AEH82071;

XX

DT 15-JUN-2007 (revised)

DT 13-JUL-2006 (first entry)

XX

DE Human gnathodiaphyseal dysplasia protein, GDD1.

XX

KW Osteopathic; Gene therapy; bone disease; bone injury; bone resorption;

KW gnathodiaphyseal dysplasia; GDD1; BOND\_PC; transmembrane protein 16E;

KW integral membrane protein GDD1; transmembrane protein 16E [Homo sapiens];

KW TMEM16E; GDD1; integral membrane protein GDD1 [Homo sapiens]; G05783;

KW G016020; G016021.

XX

OS Homo sapiens.

XX

FH Key Location/Qualifiers

FT Inhibitory-site 356

FT /note= "Missense mutations in the coding sequence can

FT lead to substitution of this residue with either Arg or

FT Gly"

XX

PN JP2006121961-A.

XX

PD 18-MAY-2006.

XX

PF 28-OCT-2004; 2004JP-00313511.

XX

PR 28-OCT-2004; 2004JP-00313511.

XX

PA (UYTO-) UNIV TOKUSHIMA NAT UNIV CORP.

XX

PI Itakura M, Tsutsumi S, Kamata N, Inoue H;

XX

DR WPI; 2006-367194/38.

DR N-PSDB; AEH82070.

DR PC:NCBI; gi47106048.

DR PC:SWISSPROT; Q75V66.

XX

PT Novel gnathodiaphyseal dysplasia DNA, useful as diagnostic agent for bone

PT disease such as gnathodiaphyseal dysplasia, bone deficiency or bone-

PT resorption property disease.

XX

PS Claim 9; SEQ ID NO 2; 11pp; Japanese.

XX

CC Revised record issued on 15-JUN-2007 : Enhanced with precomputed  
CC information from BOND.

XX  
SQ Sequence 913 AA;

Query Match 34.3%; Score 1412.5; DB 11; Length 913;  
Best Local Similarity 39.2%; Pred. No. 1.2e-141;  
Matches 309; Conservative 148; Mismatches 259; Indels 73; Gaps 20;

Qy	1	QQDVQDGNNTTVHYALLSASWAVLCYYAEDRLRLKLPQE--LPNQASNWSAGLLAWLGIPN	58
		: :   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :	
Db	120	KRDSDEGRT--YFVKIHAPWEVLVLTVAEVLGIKMPIKESDIRPKHPTISYVLGVPRLP-	176

Qy	59	VLLVVPDPVPPEYYSCRFVNKLPRFLGSDNQDTFFTTSTKRHQILFEILAKTPYGHEK-K	117
		: : :   :           : : :   : :	
Db	177	-LSVKYPH--PEYFTAFSRHROELFLIED-QATFFPSSSRNIRIVYLISRCFPGIEDGK	232

Qy 118 NLLGIQLLAEGLVLSAAFPLHDGFFKTPPEGFQAPRLNQRQVLFGHWRWGWKNKYQPLD 177  
|| :|| |:||||| : | || :| |:|||| : |||

Db 233 KRFGIERLLNSNTYSAYPLHDGOYWKPSFPPNP--TNERYLTHLNWARFSYFYKEOPLD 290

Qy	178	HVRRRYFGEKVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFSDIPTQELCGSK--DSF	235
		: : :       : :   :       :       :   : : :   :	
Db	291	LIIKKYYGKIGIYIFVLGFYTEMLFFAAVGLACFIYGLLSMEHTSIEICDPEIGGOM	350

Qy	236	EMCPLCLD-CPFWLLSSACALAQAGRLFDHGGTVVFFSLFMALWAVLLEYYWKRKSATLAY	294
		: :  : :  :  :    :    ::    :   : :  :	
Dh	351	IMCPLCDVOVCYWR NSTCLASFESHLEDNESTVFFAEMGIWVTLFLEFWMKORORAE	411

Qy	295	RWDCSIYEDTEE--RPRPQFAASAPMTAPNPITGEDEPYFPERSRARRMLAGSVVIVVMV	352
		: : ::   : :  :           :  :	
Db	411	EWDI.VDFEEEOOI.OIRFEFEAMCKHRKI.NAVTKEMEPMPI.YTRIPWYFISGATVIWM	470

Qy 353 AVVVMCLVSIILYRAIMAIIVSVSRGNTLLAAWASRI-----ASLTGS 394  
::: ::::: ::: | :::  
Db 471 SLVVTSMVAIVYVRL-----SVEATFASFMESDASIKOVKSELTPOITTSLTGS 519

**Qy**                395 VVNLVF ILILSKIYVS LAHLVLRWEMHRTQT KFDAFTLKVFIF QFVNFYSSPVYIAFFK     454  
                     : | : ||| : | : : |: | : | : | : | : | : | : | : | : | : |  
**Dd**                520 CLNFIVILT LNFEEYEKISAWIT KMFIPTRYOEYSSEI.LKMFLFOFPNVFEYSSCFYVAEFF     579

Qy 455 GRFVGYPGNYHTLFGV-RNEECAAGGCLIELAQELLVIMVGKQVINNMQEVLIPLKLGWW 513  
 |:||||| | | | :||| ||||| :| :|| ||| :|:| :| ||  
 Db 580 GKFLVGYPGKYTYLFNEWRSSECDPGCLIELTTQLTIIMTGKQIFGNIKEAIYPLALNWW 639

Qy 514 QKFLRLSKRRKAGASAGASQGPWEDDYELVPCE--GLFDEYLEMVLQFGFVTIFVAACPL 571  
 ::||| :: || |:| ||| ||| | |||||:|:| ||  
 Db 640 -----RRRKARTNSEKLYSRWEQDHDLESFGLPLGLFYEYLETVTQFGFVTLFVASFPL 692

Qy 572 APLFALLNNWVEIRLDARKFVCEYRRPVAERAQDIGIWFHILAGLTHLAVISNAFLAFLS 631  
 ||| |:|:| ||||:| | :||| || :| ||:| || |: |:| :||:|:|:  
 Db 693 APLLALINNIVEIRVDAWKLTQYRRTVASKAHSIGVWQDILYGMVLSVATNAFIVAF 752

Qy 632 SDFLPRAYRW----TRAHDLRGFLN----FTLARAPSSFAAAHNR---TCRYRAFR-- 677  
 || :|| | : :||| | :| | : | : | |||| :|  
 Db 753 SDIIPRLVYYYAYSTNATQPMGTGVNNSLSVFLIADFPNHTAPSEKRD FITCRYRDYRYP 812

Qy 678 -DDGDHY--SQTYWNLLAIRLAFVIVFEHVVSFVGRLLDLLVPDIPEVSVEIKVKREYYLA 734  
 ||: | : :|:|:| : :|:| |||| | || :|:|:| : :|:| :  
 Db 813 PDDENKYFHNMQFHWVLAAKMTFIIVMEHVFLVKFLLAWMIPDVPKDVVERIKREKLMT 872

Qy 735 KQALAENEV 743  
 : | : | :  
 Db 873 IKILHDFEL 881

## RESULT 15

ABB62812

ID ABB62812 standard; protein; 1219 AA.

XX

AC ABB62812;

XX

DT 15-JUN-2007 (revised)

DT 26-MAR-2002 (first entry)

XX

DE Drosophila melanogaster polypeptide SEQ ID NO 15228.

XX

KW Drosophila; developmental biology; cell signalling; insecticide;

KW pharmaceutical; BOND\_PC; CG6938-PA; CG6938-PA [Drosophila melanogaster];

KW CG6938.

XX

OS Drosophila melanogaster.

XX

PN WO200171042-A2.

XX

PD 27-SEP-2001.

XX

PF 23-MAR-2001; 2001WO-US009231.

XX

PR 23-MAR-2000; 2000US-0191637P.

PR 11-JUL-2000; 2000US-00614150.

XX

PA (PEKE ) PE CORP NY.

XX

PI Venter JC, Adams M, Li PWD, Myers EW;

XX

DR WPI; 2001-656860/75.

DR N-PSDB; ABL06915.

DR PC:NCBI; gi24663059.

XX

PT New isolated nucleic acid detection reagent for detecting 1000 or more  
PT genes from Drosophila and for elucidating cell signaling and cell-cell  
PT interactions.

XX

PS Disclosure; SEQ ID NO 15228; 21pp + Sequence Listing; English.

XX

CC The invention relates to an isolated nucleic acid detection reagent  
CC capable of detecting 1000 or more genes from Drosophila. The invention is  
CC useful in developmental biology and in elucidating cell signalling and  
CC cell-cell interactions in higher eukaryotes for the development of  
CC insecticides, therapeutics and pharmaceutical drugs. The invention  
CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
CC sequences (ABL01840-ABL16175) and the encoded proteins (ABB57737-  
CC ABB72072). The sequence data for this patent did not form part of the  
CC printed specification, but was obtained in electronic format directly  
CC from WIPO at ftp.wipo.int/pub/published\_pct\_sequences

CC

CC Revised record issued on 15-JUN-2007 : Enhanced with precomputed  
CC information from BOND.

XX

SQ Sequence 1219 AA;

Query Match 33.4%; Score 1378.5; DB 4; Length 1219;  
Best Local Similarity 37.8%; Pred. No. 8.5e-138;  
Matches 304; Conservative 146; Mismatches 284; Indels 71; Gaps 19;

Qy 5 QDGNITTVHYALLSASWAVLCYYAEDRLRLKP-----LQELPNQASN 45  
| | : : | : || : ||| | : :  
Db 384 QSFNEKTFFLKIHLPLWRLETRLAEVMNLKLPVKRFITISVKPSWDEENVVLRNMQYKWDV 443  
Qy 46 WSAGLLAWLGIPNVLLVVPDVPPEYYSCRFRVKNLPRFLGSDNQDTFFTSTKRHQILFE 105  
| | : : ||| : : | : : | ||| :| : :  
Db 444 WQR-LTKKIQLDQTLLE---GETTFKAATANGNPPEEQFIVKD-RATAFTSAQRSLMVMQ 497  
Qy 106 ILAKTPYGHEKKNLLGIHQLLAEGVLSAAFFLHDGFPFKTPPEGPQAPRLN-QRQVLFQHW 164  
:| :|| : : || : : :| |||| :| : : : : :| :| :|  
Db 498 VLIRTPFDESQRS--GIRRLMNDGTYLGCFFLHEGRY---DRPHSSGISLDRRVLYQTW 551



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Qy      165 ARWGWKNKYQPLDHRVRYFGEKVALYFAWLGFYTGWLLPAAVVGTLVFLVGCFLVFS-- 222
      | :| | ||| ||:||||:|:|||| |||| | : | |||| | : | : | :
Db      552 AHPSQWYKQPLCLVRKYFGDKIALYFCWLGFYTEMLVYPVAVGTLCFIYGLATLESEDN 611

Qy      223 IPTQELCG--SKDSFEMCPLC-LDCPFWLLSSACALAQAGRLFDHGGTVFFSLFMALWAV 279
      |::|:| : : |||| | : || :| :: |||: |||::||: ||
Db      612 TPSKEICNEYGTGNITLCPCLKACSYQRLSESCLSRLTYLFDNPNSTVFFFAIFMSEFWAT 671

Qy      280 LLELYWKRKSATLAYRWDCSDYEDTEERPRPQFAASAPMTAPNPITGEDEPYFPERSRAR 339
      || |||| : | : || : | :| ||:| :| ||:| | ||| :|:
Db      672 TFLELWKRKQSVLVWEWDLHNV-DMDEENRPEFETNATTFRMNPVTREKEPYMSTWNRSI 730

Qy      340 RMLAGSVVIVVMVAVVMCLVSIILYRAIMAIVVSRSGNTLLAAWASRIASLTGSVVNLV 399
      | : : : ||::|: : ||| : | : | : | :| :| :| :|
Db      731 RFVITGSAVLFMISVVL SAVLGTILYRITLVSVIYGGGGFFVKEHAKLFTSVTAALINLV 790

Qy      400 FILILSKIYVSLAHLVLRWEMHRTQTKFEDAFTLKVFIFQFVNFYSSPVYIAFFKGRFVG 459
      |::|::|| :| || | || ||::||:| || :|::| ||| :| |||||
Db      791 VIMILTRIYHRMAIKLTNLENPRTHTEYEDSYTFKIFFFEFNMFYSSLIYIAFFKGRFFD 850

Qy      460 YPGNYHT---LFGVRNEECAAGGCLIELAQELLVIMVGKQVINMQEVLIPKLKGWQWK 515
      |||: |::| :|| : | || || :|: |||| | | || | | :|
Db      851 YPGDDQARKSEFFRLKNDICDPAGCLSELCLQLAIIIMVGKQCWNFMFYLPKFWNWNR- 909

Qy      516 FRLRSKKRKAGASAGASQGPWEDDYELV-PCE-GLFDEYLEMVLQFGFVTIFVAACPLAP 573
      : | ::| || || : | ||||| ||: ||||| ||| ||| |||
Db      910 ---QRKHQATKDESHLMAWEQDYHMQDPGRLLALFDEYLEMILQYGFVTLFVAAPFLAP 966

Qy      574 LFALLNNWEIRLDARKEVCEYRRPVAERAQDIGIWFHILAGLTHLAVISNAFLAFSSD 633
      ||||| ||||| | : ||: ||| :|| : | :| :|: ||||: ||: ||
Db      967 LFALLNNVAEIRLDAYKMVTQARRPLAERVEDIGAWYGILRIITYTAVVSNAFVIATSD 1026

Qy      634 FLPRAYYR--WTRAHDLRGLNFTLA-----RAPSSFAAAHNRTCRYRAFRDDGDH 682
      |::| | : : | | :| :| :| :| :| :| :| :| :| :| :|
Db      1027 FIPRMVYKFVYSETHTLAGYIEHSLSIFNTSDYKEEWGASVSEKDPDTCQYRGYRNGPKD 1086

Qy      683 Y-----SQTYWNLLAIRLAFVIVFEHVVSFVGRLLDLLVPDIPESVEIKVKREYYLAKQA 737
      | | | :| ||||: ||||| : : :||:| | : :||: |||:|
Db      1087 YEPYGLSPHYWHFAARLAFVVVFEHVVFITGIMQFIIPDVPSEVKTQMQRQLLAKEA 1146

Qy      738 LAENEVLFGTNGTKDEQPKGSELSS 762
      : :| | | :| :|
Db      1147 KYQ-----HGIKRAQGDSQDIMS 1164

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Search completed: June 24, 2008, 15:22:07

Job time : 273 secs

SCORE 0.0